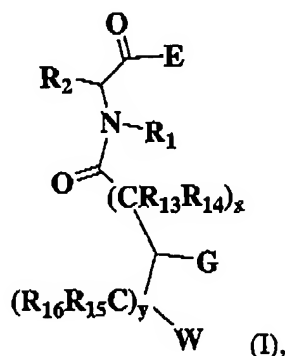


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of claims:

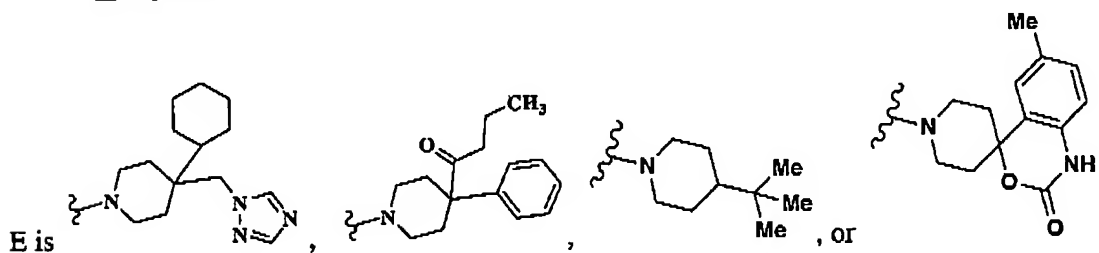
1. (Currently amended) A compound of formula (I),



or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

R_1 is hydrogen or C_{1-6} alkyl or is taken together with R_2 or R_3 to form a monocyclic or bicyclic aryl, cycloalkyl, heteroaryl or heterocycle;

R_2 is C_{1-6} alkyl or C_{2-6} alkenyl optionally substituted with one to three aryl, cycloalkyl, or heteroaryl, provided that where G is C_{2-6} alkenyl, $A_4-NR_{18}CO_2R_{19}$, or $A_4-SO_2R_{17}$, or when y is 0, R_2 ~~is may be~~ or C_{1-6} alkyl or C_{2-6} alkenyl, each optionally substituted with heteroaryl;



G is selected from $A_4-NR_{18}C(=O)R_{19}$, $A_4-NR_{18}SO_2R_{17}$, $A_4-NR_{18}CO_2R_{19}$, and

$A_4-NR_{20}C(=O)NR_{18}R_{19}$ wherein A_4 is a bond, C_{1-6} alkylene, or C_{2-6} alkenylene, or where G is $A_4-NR_{18}CO_2R_{19}$, or when y is 0, R_2 ~~is may be~~ C_{1-6} alkyl or C_{2-6} alkenyl, each optionally substituted with heteroaryl;

W is selected from substituted or unsubstituted heterocyclo, heteroaryl, or cycloalkyl selected from azetidiny and imidazolyl, each optionally substituted with lower alkyl;

~~R₁₃, R₁₄, R₁₅ and R₁₆ are hydrogen-selected independently of each other from hydrogen, alkyl, substituted alkyl, amino, alkylamino, hydroxy, alkoxy, aryl, cycloalkyl, heteroaryl, or heterocyclo, or R₁₃ and R₁₄, or R₁₅ and R₁₆, when attached to the same carbon atom, may join to form a spirocycloalkyl ring;~~

R₁₇ is alkyl, substituted alkyl, cycloalkyl, aryl, heterocyclo, or heteroaryl;

R₁₈, R₁₉, and R₂₀ are independently selected from hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, aryl, heteroaryl, cycloalkyl, heterocyclo, or C(=O)R₂₈; or when G is NH(C=O)R₁₉, R₁₉ may be a bond joined to W to define a heterocyclo ring; provided, however, that when y is at least one, W is imidazolyl, indolyl, ~~NR₂₁R₂₂, or OR₂₃~~, and G is ~~NR₁₈C(=O)R₁₉~~, then R₁₉ is not a C₁-alkyl having the substituent ~~NR₂₉R₃₁~~;

R₂₉ and R₃₁ are selected from hydrogen, alkyl, haloalkyl, hydroxyalkyl, phenylalkyl, and alkoxyalkyl, or R₂₉ and R₃₁ taken together form a heterocyclo ring;

x is 0, 1, or 2; and

y is 0, 1, 2, 3 or 4.

2. (Currently amended) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

G is selected from:

a) ~~NR₁₈C(=O)R₁₉~~;

b) ~~C₁₋₆alkylene or C₂₋₆alkenylene joined to one of NR₁₈C(=O)R₁₉, NR₁₈CO₂R₁₉, NR₁₈SO₂R₁₇, and NR₂₀C(=O)NR₁₈R₁₉~~;

R₁₇ is C₁₋₄alkyl, C₅₋₆cycloalkyl, phenyl, or benzyl;

R₁₈, R₁₉, and R₂₀ are independently selected from hydrogen, C₁₋₄alkyl, phenyl, benzyl, C₅₋₆cycloalkyl, -C(=O)CH₂(phenyloxy), -C(=O)CH₂(benzyloxy), imidazolyl, pyridyl, furyl, thienyl, or C₁₋₄alkyl or C₂₋₄alkenyl substituted with one of phenyl, pyridyl, furyl, cyclopentyl, cyclohexyl, CO₂Me, phenyloxy, or benzyloxy, wherein each ringed group of R₁₈, R₁₉, and R₂₀ in turn is optionally substituted with one to two R₃₆, and/or optionally has a benzene ring or five membered heterocyclo having two oxygen atoms fused thereto; and

R_{36} is halogen, methoxy, nitro, phenyl, phenyloxy, or alkylamino.

3. (Currently amended) A compound according to claim 2, or a pharmaceutically-acceptable salt or hydrate, thereof, in which

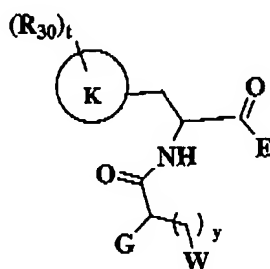
G is $\text{NR}_{18}\text{C}(=\text{O})\text{R}_{19}$,

R_{18} is hydrogen or lower alkyl, and

R_{19} is C_{1-4} alkyl, C_{2-4} alkenyl, phenyl, benzyl, C_{5-6} cycloalkyl, $-\text{C}(=\text{O})\text{CH}_2(\text{phenyloxy})$, $-\text{C}(=\text{O})\text{CH}_2(\text{benzyloxy})$, imidazolyl, pyridyl, furyl, thienyl, or C_{1-4} alkyl or C_{2-4} alkenyl substituted with one of phenyl, phenyl, pyridyl, furyl, cyclopentyl, cyclohexyl, CO_2Me , phenyloxy, and benzyloxy, wherein each ringed group of R_{19} in turn is optionally substituted with one to two R_{36} , and/or optionally has a benzene ring or five membered heterocycle having two oxygen atoms fused thereto.

4. (Previously Presented) A compound according to claim 2, or a pharmaceutically-acceptable salt or hydrate, thereof, in which W is azetidiny or imidazolyl.

5. (Previously Presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, having the formula:



in which

K is phenyl or thiazolyl;

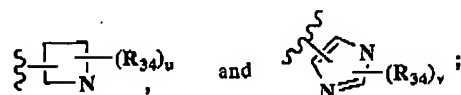
R_{30} is selected from C_{1-4} alkyl, hydroxy, alkoxy, halogen, nitro, cyano, amino, alkylamino, phenyl, and $-\text{C}(=\text{O})\text{phenyl}$;

t is 0, 1 or 2; and

y is 0, 1 or 2.

6. (Canceled)

7. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which
W is a ring selected from:



R₃₄ at each occurrence is attached to any available carbon or nitrogen atom of W and is selected from

C₁₋₆alkyl

u is selected from 0, 1, 2, and 3; and

v is 0, 1 or 2.

8. - 9. (Canceled)

10. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which

R₂ is selected from C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkenylene-K, and -(CH₂)_g-K;

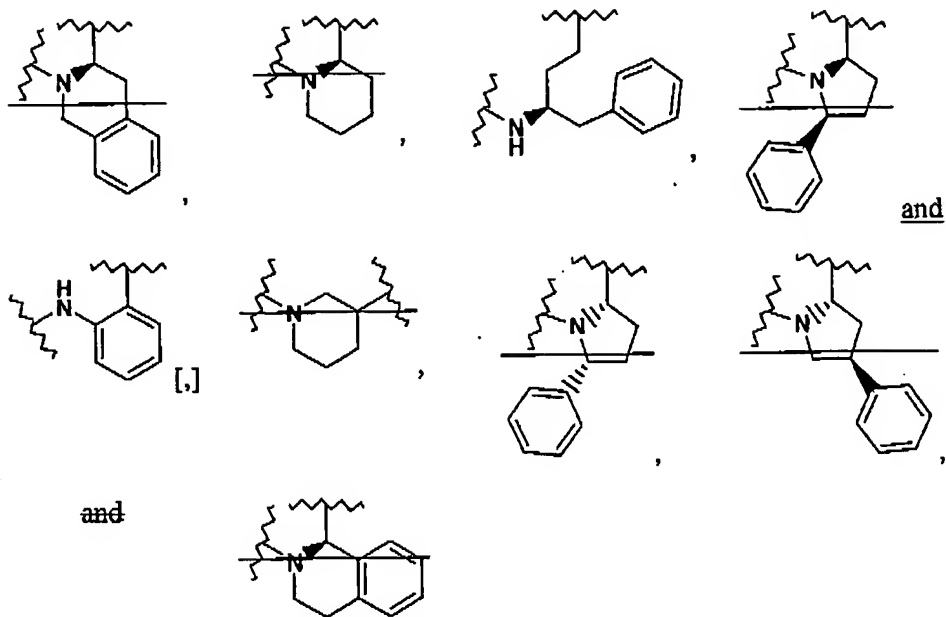
K is selected from phenyl, naphthyl, thienyl, thiazolyl, pyridinyl, pyrimidinyl, and C₅₋₆cycloalkyl,

wherein each group K in turn is optionally substituted with one to three R₃₀ or has a benzene ring fused thereto, which also may be substituted with one to three R₃₀;

R₃₀ is selected from C₁₋₄alkyl, hydroxy, alkoxy, halogen, nitro, cyano, amino, alkylamino, phenyl, and acylphenyl; and

g is 0, 1, 2 or 3.

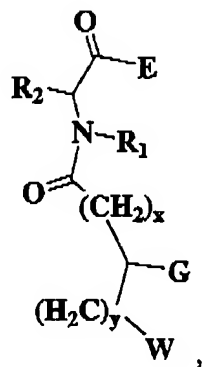
11. (Currently Amended) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which -N(R₁)-CH(R₂)- taken together are selected from,



12. (Previously presented) A compound according to claim 1, or a pharmaceutically-acceptable salt or hydrate, thereof, in which R_1 is hydrogen or C_{1-4} alkyl.

13. (Canceled)

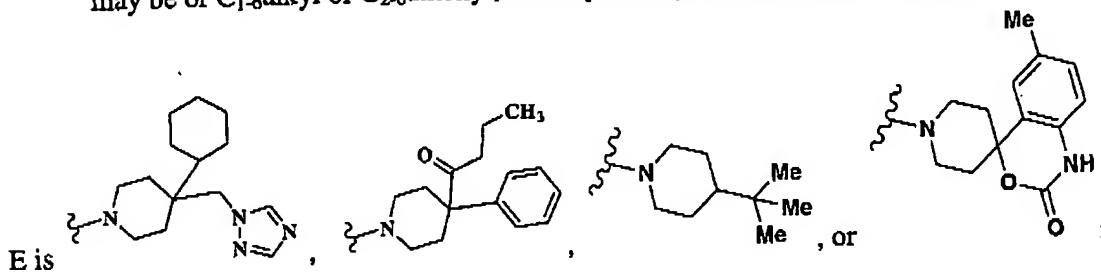
14. (Currently amended) A compound having the formula,



or a pharmaceutically-acceptable salt or hydrate, thereof, in which:

R_1 is hydrogen or C_{1-6} alkyl or is taken together with R_2 or R_3 to form a monocyclic or bicyclic aryl, cycloalkyl, heteroaryl or heterocycle;

R_2 is C_{1-6} alkyl or C_{2-6} alkenyl optionally substituted with one to three aryl, cycloalkyl, or heteroaryl, provided that where G is C_{2-6} alkenyl, or $[A_1]-NR_{18}CO_2R_{19}$, or $A_1-SO_2R_{17}$ or when y is 0, R_2 may be or C_{1-6} alkyl or C_{2-6} alkenyl, each optionally substituted with heteroaryl;



G is selected from:

a) $NR_{18}C(=O)R_{19}$;

b) C_{1-6} alkylene or C_{2-6} alkenylene joined to one of $NR_{18}C(=O)R_{19}$, $NR_{18}CO_2R_{19}$, $NR_{18}SO_2R_{17}$, and $NR_{20}C(=O)NR_{18}R_{19}$;

W is selected from -- substituted or unsubstituted heterocyclo, heteroaryl, or cycloalkyl selected from azetidiny and imidazolyl, each optionally substituted with lower alkyl;

R_{17} is alkyl, substituted alkyl, cycloalkyl, aryl, heterocyclo, or heteroaryl;

R_{18} , R_{19} , and R_{20} are independently selected from hydrogen, alkyl, alkenyl, aryl, heteroaryl, cycloalkyl, heterocyclo, $C(=O)R_{28}$ or a C_{1-4} alkyl or C_{2-4} alkenyl substituted with one or more of aryl, heteroaryl, cycloalkyl, heterocyclo, alkoxycarbonyl, phenyloxy, and/or benzyloxy, and each of said ringed groups of R_{18} , R_{19} , and R_{20} in turn is optionally substituted with one to two R_{36} ;

R_{21} and R_{22} are selected from alkyl and substituted alkyl;

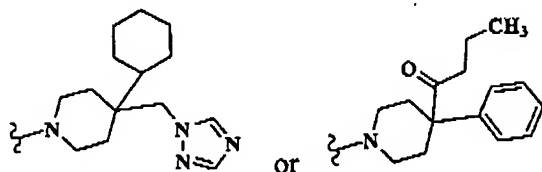
R_{36} is halogen, methoxy, nitro, phenyl, phenyloxy, or alkylamino;

x is 0, 1, or 2; and

y is 0, 1, 2, 3 or 4.

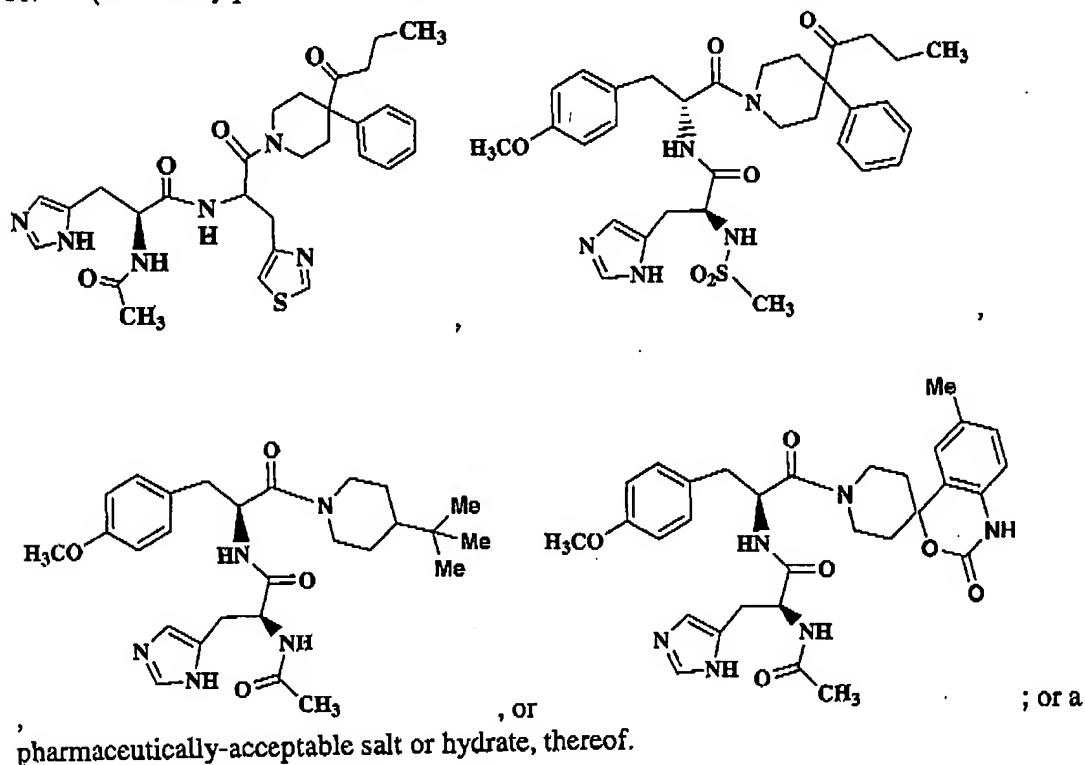
15. (Canceled)

16. (Previously presented) A compound according to claim 14, or a pharmaceutically-acceptable salt or hydrate, thereof, in which E is



17. (Previously presented) A compound according to claim 14, or a pharmaceutically-acceptable salt or hydrate, thereof, in which G is $\text{NHC}(=\text{O})(\text{alkyl})$ or $\text{NHC}(=\text{O})\text{phenyl}$.

18. (Previously presented) A compound according to claim 1, having the formula,



19. (Previously presented) A pharmaceutical composition comprising at least one compound according to claim 1 or a pharmaceutically-acceptable salt or hydrate, thereof; and a pharmaceutically-acceptable carrier or diluent.

20. - 23. (Canceled)